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POSITION STATEMENT

Conjunctivitis in atopic dermatitis patients with and without dupilumab therapy – international eczema council survey and opinion

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Abstract

Background Conjunctivitis is common in patients with atopic dermatitis (AD) in general and a commonly reported adverse event in AD clinical trials with dupilumab.

Objective To survey opinions and experience about conjunctivitis occurring in AD, including those during dupilumab treatment in a group of AD experts from the International Eczema Council (IEC).

Methods Electronic survey and in-person discussion of management strategies.

Results Forty-six (53.5%) IEC members from 19 countries responded to the survey. Consensus was reached for several statements regarding diagnostic workup, referral and treatment. IEC members suggest that patients with AD should (i) routinely be asked about ocular complaints or symptoms, (ii) obtain information about the potential for conjunctivitis before starting dupilumab therapy and (iii) if indicated, be treated with dupilumab despite previous or current conjunctivitis. In cases of new-onset conjunctivitis, there was consensus that dupilumab treatment should be continued when possible, with appropriate referral to an ophthalmologist.

Limitations The study relies on expert opinion from dermatologists. Responses from few dermatologists without dupilumab access were not excluded from the survey.

†Correction added on 29 June 2019, after first online publication: the surname of author 'C. Traidl-Hoffmann' has been corrected.

Conclusion The IEC recommends that dermatologists address conjunctivitis in patients with AD, especially during treatment with dupilumab.

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Conflict of interest

Dr. Thyssen has attended advisory boards for Roche, Eli Lilly and Sanofi Genzyme and received speaker honorarium from LEO Pharma and Sanofi Genzyme. Dr. de Bruin-Weller M has been a principal investigator, advisory board member and consultant for Regeneron Pharmaceuticals, Inc. and Sanofi Genzyme and a principal investigator, advisory board member for AbbVie. Dr. Paller has been a principal investigator for AbbVie, Anaptysbio, Eli Lilly, Galderma, Incyte, Leo, Janssen, Novartis and Regeneron, and a consultant for AbbVie, Amgen, Asana, Dermavant, Dermira, Galderma, Eli Lilly, Forte, Leo, MatriSys, Menlo, MorphoSys/Galapagos, Novartis, Pfizer, Regeneron and Sanofi. Dr. Leshem has been a principal investigator, consultant and/or advisory board member for and/or received honoraria from AbbVie Inc., Sanofi and Regeneron Pharmaceuticals Inc., Pfizer, Dexcel Pharma, Genentech and Eli Lilly. Dr. Vestergaard has been a investigator, speaker or consultant for Novartis, AbbVie, Sanofi, Leo Pharma and Eli Lilly. Dr. Deleuran has been a principal investigator, speaker, advisory board member and/or consultant for LEO Pharma, Lilly, Novartis, Pfizer, Regeneron Pharmaceuticals, Inc., Sanofi Genzyme, MorphoSys-Galapagos, Pierre Fabre and MEDA. Dr. Drucker has served as a investigator and has received research funding from Sanofi and Regeneron and has been a consultant for Sanofi, RTI Health Solutions, Eczema Society of Canada and Canadian Agency for Drugs and Technology in Health. He has received honoraria from Prime Inc, Spire Learning, CME Outfitters and Eczema Society of Canada. His institution has received educational grants from Sanofi. Dr. Fölster-Holst has been a investigator, speaker or consultant for Astellas, Almirall Hermal, Beiersdorf, Johnson & Johnson, La Roche-Posay, LEO Pharma, Neubourg GmbH, Novartis, Pierre Fabre, Procter & Gamble and Regeneron Pharmaceuticals. Dr. Traidl-Hoffmann has been a investigator, speaker or consultant for Novartis, Lilly, Regeneron Pharmaceuticals, Sanofi Genzyme and Sebapharma. Dr. Kilian Eyerich has been a principal investigator, advisory board member or consultant for AbbVie, Almirall, Berlin-Chemie, Hexal, Janssen, Leo, Lilly, Novartis and Sanofi. Dr. Alain Taieb has been a consultant or investigator for Pierre Fabre, Galderma, Novartis, Johnson and Johnson and Sanofi. Dr. Su has been a principal investigator, consultant or advisory board member for Amgen, AbbVie, Eli Lilly, Janssen, Meda, Novartis, Pfizer, Pierre Fabre and Sanofi. Dr. Bieber has been a principal investigator, advisory board member or consultant for Regeneron, Sanofi, GSK, Celgene, AbbVie, AnaptysBio, MedImmune, Chugai, Pierre Fabre, Novartis, Asana Biosciences, LEO, Galapagos/MorphoSys, BioVersys, Galderma, Kymab, Glenmark, Astellas, Daiichi-Sankyo, Lilly, Pfizer, MenloTx, Dermavant and Almirall. Dr Michael J. Cork is a investigator and consultant for Regeneron, Sanofi Genzyme, Pfizer, Leo, Galapagos, Novartis, Boots, L'Oreal, Dermavant, Menlo, Reckitt Benckiser, Oxagen, Johnson & Johnson, Hyphens, Astellas, AbbVie, Galderma and Procter & Gamble. Dr. Eichenfield has been a principal investigator, consultant or advisory board member for Amgen, Dermavant, Dermira, Eli Lilly, Forte, Galderma, Leo, MatriSys, Menlo, MorphoSys, Novartis, Ortho Dermatologics, Pfizer, Pierre Fabre and Regeneron/Sanofi. Dr Guttman-Yassky is an employee of Mount Sinai and has received research funds (grants paid to the institution) from AbbVie, Celgene, Eli Lilly, Janssen, MedImmune/Astra Zeneca, Novartis, Pfizer, Regeneron, Vitae, Glenmark, Galderma, Asana, Innovaderm, Dermira, UCB, Kiniksa and Novan. EGY is also a consultant for Sanofi Aventis, Regeneron, Stiefel/GlaxoSmithKline, MedImmune, Celgene, AnaptysBio, Dermira, Galderma, Glenmark, Novartis, Pfizer, Vitae, Leo Pharma, AbbVie, Eli Lilly, Kyowa Kirin, Mitsubishi Tanabe, Asana Biosciences, Promius, Kiniksa, Novan and Evelo biosciences. Dr. Wollenberg has been a principal investigator, advisory board member or consultant for Galderma, LEO Pharma, Lilly, MedImmune, Novartis, Pfizer, Regeneron Pharmaceuticals, Inc. and Sanofi Genzyme, and received speaker honoraria from Chugai, Galderma, LEO Pharma, MedImmune, Pfizer, Regeneron Pharmaceuticals, Inc. and Sanofi Genzyme. Corporate sponsorship was provided to the International Eczema Council by AbbVie, Asana, Celgene, Chugai, Dermavant, Dermira, Eli Lilly, Galderma, Leo Pharma, Novartis, Pfizer, Pierre Fabre, Sanofi, Genzyme and Regeneron Pharmaceuticals, Sienna and Valeant. This research was performed independently through the authors' academic university and hospital affiliations.

Introduction

Atopic dermatitis (AD) is a chronic inflammatory skin condition.¹ Patients with AD have an increased risk of developing keratoconus, cataract, glaucoma and blepharitis, as well as allergic, atopic, vernal and infectious (kerato) conjunctivitis.^{2,3}

Clinical trials with dupilumab, a monoclonal antibody that inhibits interleukin (IL)-4 and IL-13 signalling, have demonstrated efficacy in moderate-to-severe AD. However, trials have also shown an increased incidence of conjunctivitis with dupilumab treatment,^{4–6} which in some cases leads to cessation of treatment.⁷ The clinical features of dupilumab-associated conjunctivitis include bilateral inflammation of the anterior conjunctiva and hyperaemia of the limbus, which may cause nodular swelling.⁸ In a case series, dupilumab-associated conjunctivitis appeared to occur more often in patients with severe AD at baseline and with atopic comorbidities. At present, however, predictive factors are unknown.⁷

There is no literature-based guidance available about management of dupilumab-associated conjunctivitis. A group of AD experts, all councillors or associates of the International Eczema Council (IEC), therefore addressed this issue by completing a survey and discussing conjunctivitis and its management in patients with AD.

Materials and methods

The IEC consists of 86 AD experts from 22 countries (<http://www.eczemacouncil.org/>). In May 2018, a survey developed by three of the authors (J.P.T, M.S.d.B.W. and A.W.) was sent electronically to IEC councillors and associates to examine their opinions regarding management of conjunctivitis in patients with AD focusing on dupilumab-associated conjunctivitis. The survey consisted of questions and statements followed by possible responses of 'strongly disagree', 'disagree', 'neither agree, nor disagree', 'agree' and 'strongly agree' as well as closed response categories (for detailed questions, Table 1). All responses were anonymous. No *a priori* definition of consensus was determined but, post hoc, and it was decided to use rules similar to the Harmonizing Outcome Measures for Eczema (HOME) initiative, which have been used in previous IEC consensus articles.^{9,10} Consensus required that less than 30% of the voters disagreed (i.e. no more than 30% had to mark the following responses 'strongly disagreed' or 'disagreed') for each question with a response rate of at least 90% of respondents in the survey. Descriptive data analysis was performed using Excel (Windows). On 13 September 2018, a round table discussion among IEC members was held on the topic at the European Academy of Dermatology and Venereology meeting in Paris, France. In the remainder of this manuscript, IEC members who completed the survey are respondents; IEC members who participated in the discussion participants.

Results

Survey results

The following sections highlight key findings from the survey. Question and response details are shown in Table 1.

Respondent characteristics

A total of 46 councillors and associates from six continents responded to the survey (participation rate 53.5%). Respondents came from Germany ($n = 8$), the United States ($n = 7$), the Netherlands ($n = 4$), United Kingdom ($n = 4$), Denmark ($n = 3$), France ($n = 3$), Korea ($n = 2$), Japan ($n = 2$), Australia ($n = 2$) and Canada ($n = 2$), as well as, Brazil, China, India, Ireland, Israel, Italy, Switzerland, Taiwan and Tanzania (each $n = 1$).

A total of 38% and 22% of respondents reported seeing 0–20 paediatric and adult patients with AD per week, respectively, whereas 27% and 31% of respondents reported seeing 21–40 paediatric and adults patients with AD per week, respectively. The proportion of respondents that reported seeing 41–100 paediatric and adult patients per week was 31% and 33%, respectively. Remaining respondents reported seeing more than 100 paediatric and adult patients with AD per week.

Dermatologist-estimated frequency of ocular surface disease and diagnostic workup

Respondents reported relatively frequent occurrence of conjunctivitis and blepharitis in their patients with AD (Fig. 1a and b). As an indication of disease severity, half of the respondents reported having seen a paediatric or adult AD patient who required corneal transplant. Consensus was reached that patients with AD routinely should be asked about ocular complaints or symptoms, and in case of conjunctivitis, routinely referred to an ophthalmologist for diagnostic workup and treatment. Consensus was reached that AD patients with conjunctivitis should undergo (or be referred to) skin prick or specific IgE testing for common aeroallergen and be referred for patch testing with an ophthalmological series and native eye drops and ointments to detect possible delayed-type hypersensitivity to topical ingredients.

Treatment

Consensus was reached that dermatologists should initiate relevant treatment with eye drops/ointment, or oral antihistamines, for conjunctivitis before referring to an ophthalmologist, but also dermatologists should leave the indication to use of corticosteroid, ciclosporin or tacrolimus eye drops for conjunctivitis to ophthalmologists. Consensus was reached that patients with AD should be informed about possible conjunctivitis before prescribing dupilumab. There was also consensus that previous, (kerato) conjunctivitis in patients with AD should not prevent

Table 1 Overview of responses given by IEC members in a survey about conjunctivitis. Statements that reached consensus are highlighted with grey. Consensus required that less than 30% of the voters disagreed (i.e. no more than 30% had to mark the following responses 'strongly disagreed' or 'disagreed') for each question with a response rate of at least 90% of the respondents.

Survey Questions	Strongly agree		Agree		Neither agree nor disagree		Disagree		Strongly disagree	
	%	n	%	n	%	n	%	n	%	n
Atopic dermatitis patients should routinely be asked about ocular complaints or symptoms	55.81	24	32.56	14	9.30	4	2.33	1	0.00	0
I have observed corneal transplant in one or several of my atopic dermatitis patients where I assume it was due to excessive rubbing (both children and adults)	15.91	7	34.09	15	15.91	7	22.73	10	11.36	5
Atopic dermatitis patients with conjunctivitis should routinely be referred to an ophthalmologist for diagnostic work-up and treatment	23.26	10	53.49	23	9.30	4	13.95	6	0.00	0
Dermatologists should initiate relevant treatment with eye drops/eye ointment or oral antihistamines for conjunctivitis themselves before referring to an ophthalmologist	6.82	3	40.91	18	31.82	14	18.18	8	2.27	1
Atopic dermatitis patients with conjunctivitis should undergo (or be referred to) skin prick testing (or specific IgE) for common aeroallergens	25.58	11	25.58	11	30.23	13	16.28	7	2.33	1
Atopic dermatitis patients with conjunctivitis should undergo (or be referred to) skin prick testing to eyedrops if such are used	9.09	4	25.00	11	25.00	11	38.64	17	2.27	1
Atopic dermatitis patients with conjunctivitis should undergo (or be referred to) patch testing with a standardized ophthalmologic series, and native eye drops/ointments to diagnose possible delayed type hypersensitivity reactions to topical ingredients if these are used	9.30	4	41.86	18	23.26	10	25.58	11	0.00	0
Dermatologists should leave the indication to use of ciclosporin or tacrolimus eye drops for conjunctivitis to ophthalmologists	18.18	8	36.36	16	25.00	11	18.18	8	2.27	1
Dermatologists should leave the indication to use of corticosteroid eye drops for conjunctivitis to ophthalmologists	15.91	7	38.64	17	18.18	8	25.00	11	2.27	1
It is important to routinely inform about possible conjunctivitis in your atopic dermatitis patients before prescribing dupilumab (Dupixent)	59.09	26	31.82	14	6.82	3	2.27	1	0.00	0
Patients with atopic dermatitis should be referred to an ophthalmologist before initiation of dupilumab therapy	4.88	2	14.63	6	14.63	6	56.10	23	9.76	4
The risk of conjunctivitis when using dupilumab may prevent me from using the drug in patients with previous (kerato-) conjunctivitis?	4.55	2	13.64	6	22.73	10	47.73	21	11.36	5
The risk of conjunctivitis when using dupilumab may prevent me from using the drug in patients with current (kerato-) conjunctivitis?	4.65	2	41.86	18	25.58	11	23.26	10	4.65	2
Patients with new onset conjunctivitis during dupilumab treatment should always be referred to an ophthalmologist	20.45	9	47.73	21	20.45	9	11.36	5	0.00	0
Patients with new onset conjunctivitis during dupilumab treatment should be referred to an ophthalmologist in more severe cases, (inadequate response to artificial tears and/or antihistamine eye drops)	50.00	22	34.09	15	6.82	3	9.09	4	0.00	0
New onset conjunctivitis during dupilumab should result in referral of the patient to an ophthalmologist but treatment should be continued	11.36	5	56.82	25	25.00	11	6.82	3	0.00	0
New onset conjunctivitis during dupilumab should result in referral of the patient to an ophthalmologist but treatment should be paused	4.55	2	4.55	2	31.82	14	52.27	23	6.82	3

the use of dupilumab (Fig. 2; for discussion on dupilumab treatment initiation in patients with current conjunctivitis, see detailed discussion below). There was consensus that patients with new-onset conjunctivitis during dupilumab treatment should always be referred to an ophthalmologist, but also that dupilumab treatment should be continued while awaiting ophthalmologist consultation.

Round table discussion

The survey results were presented at the meeting in Paris where 20 councillors and associates were present. Most of the

discussion concerned dupilumab-associated conjunctivitis, which most IEC members had seen and treated in their adult patients with AD. Dupilumab-associated conjunctivitis had also been seen in paediatric patients treated in clinical trials with dupilumab. It was agreed that secondary analyses of completed clinical trials with dupilumab and ongoing postmarketing registries are urgently needed to identify sub-groups with increased risk of conjunctivitis after dupilumab administration. Better understanding of the aetiopathogenesis of dupilumab-associated conjunctivitis to identify susceptible patients and future treatments was emphasized. Although conjunctivitis has been

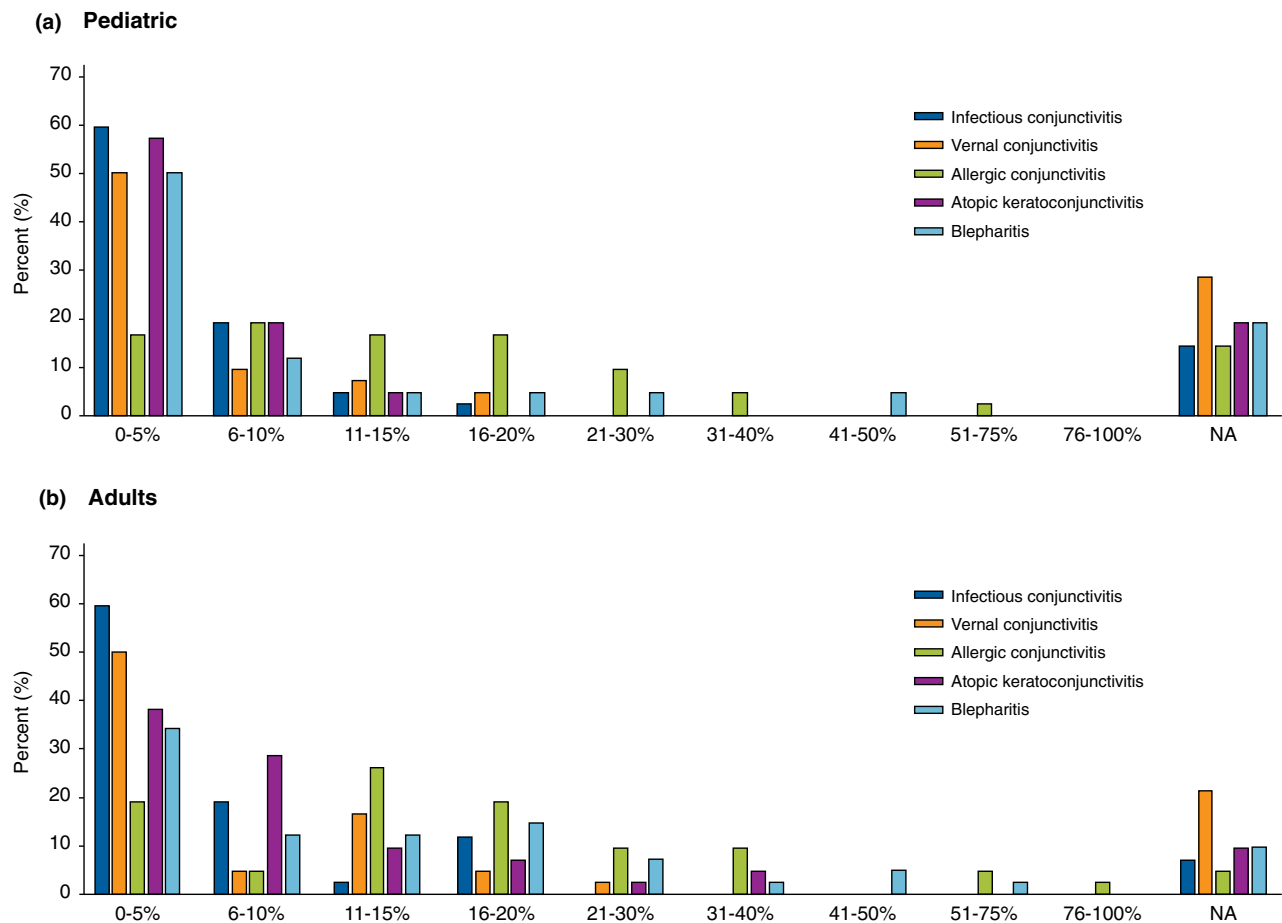


Figure 1 Proportion of IEC member who report having seen pediatric (a) and adult (b) patients with AD and who have been diagnosed with conjunctivitis and blepharitis within the 7 past 12 months.

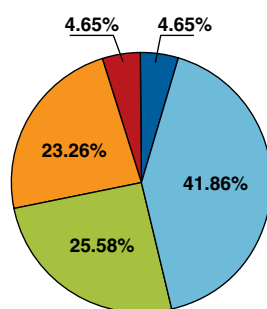
associated with the experimental use of IL-13 inhibitors for AD, the numbers in trials to date are small and discussants warned about the possibility of similar ocular complications in AD patients treated with IL-13 inhibitors.

It was the experience of IEC members that many AD patients with dupilumab-associated conjunctivitis clear their ocular symptoms, some spontaneously and some after a short course of treatment with lubricating eye drops and ointments containing corticosteroids or tacrolimus. Several IEC members had good experiences with the use of fluorometholone 0.1% eye drops or tacrolimus 0.03% ointment.⁸ Corticosteroid eye drops should be used with caution, since they increase the risk of superinfection and glaucoma. Nevertheless, few patients stop dupilumab treatment due to conjunctivitis and conjunctivitis may improve dramatically during dupilumab use. These varied observations regarding course further stress the need to study these patients and find correlates with outcome. Some IEC members recommended artificial lubricating tears when starting a patient on

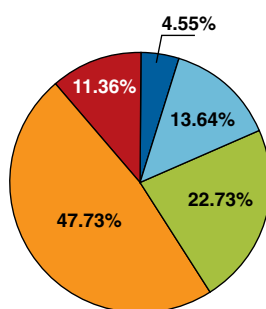
dupilumab as prevention to reduce ocular complaints and the risk of conjunctivitis but, while there is a scientific rationale for this intervention, more work is needed to develop optimal regimens and assess its efficacy. A strategy pursued by some IEC members is to taper dupilumab injections e.g. to every 3rd week, but potential correlation with extending the dosing interval and increasing the risk of developing anti-drug antibodies against dupilumab was stressed.

The clinical course and prognosis of already existing conjunctivitis following initiation of dupilumab is difficult to predict. In the experience of IEC councillors, some patients with AD may experience worsening of ocular symptoms due to unrecognized effects of dupilumab treatment, some may experience stable and unchanged conjunctivitis, and some may experience improvement and even complete resolution of ocular symptoms. Patients should be informed about the potential of conjunctivitis with dupilumab and should be referred to an ophthalmological when clinically indicated. It was stressed that at this moment

The risk of conjunctivitis when using dupilumab may prevent me from using the drug in patients with current (kerato-) conjunctivitis?



The risk of conjunctivitis when using dupilumab may prevent me from using the drug in patients with previous (kerato-) conjunctivitis?



Patients with new onset conjunctivitis during dupilumab treatment should always be referred to an ophthalmologist

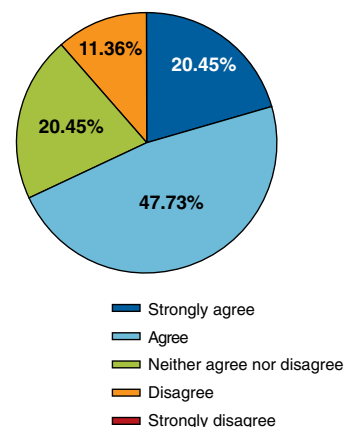


Figure 2 Key findings from the survey regarding management of AD patients with conjunctivitis treated with dupilumab.

there are also no reliable data on the course of atopic keratoconjunctivitis and vernal keratoconjunctivitis during dupilumab treatment. Patient experiencing these should be carefully monitored by an ophthalmologist before and during treatment with dupilumab. The AD severity, conjunctivitis severity, possible contraindications, possible effect of dupilumab therapy on concomitant asthma or other comorbidities, as well as other treatment options, should be considered on an individual patient basis.

Discussion

This survey of IEC councillors and associates, representing 19 countries from six continents, examined opinions regarding proper management of paediatric and adult patients with AD with conjunctivitis, and in particular, conjunctivitis occurring during dupilumab treatment. Participants in this study care for many patients with AD weekly.

Few studies have examined the exact prevalence and incidence of conjunctivitis in patients with AD. In young children from a U.S. trial, the prevalence of allergic conjunctivitis was 16%,¹¹ whereas chronic conjunctivitis and superficial keratitis were observed in 32% and 7% of 44 adult patients with AD when examined ophthalmologically.¹² In Danish adults, 12% and 18% of adults with mild and severe AD, respectively, had used anti-inflammatory ocular agents, and 13 of 1000 severe patients with AD per year had used an ocular anti-inflammatory agent compared with 6 of 1000 individuals (sex- and age-adjusted) from the general population.³ These data emphasize that conjunctivitis is more common in patients with AD than controls and, accordingly, the IEC reached consensus and recommended that patients with AD routinely should be asked about signs and

symptoms of conjunctivitis. Recommendations about eye examination for conjunctivitis in AD, as well as diagnostic workup, referral and proper management, have not been included in recent European and North American guidelines for AD.^{1,13–18}

The estimated prevalence of ocular surface disease provided by the IEC members should be interpreted with caution because of the risk of recall bias and the skewed population towards more severe disease seen by IEC members. Moreover, there was no information about the severity or accuracy of diagnosis of the conjunctivitis (e.g. if ophthalmologist-confirmed). Nonetheless, IEC members experienced that adults with AD frequently suffer from conjunctivitis and blepharitis. It is currently unclear whether blepharitis should be regarded as part of AD or, instead, occurs as a result of co-pathogenic factors and is a complication or comorbidity. Importantly, other aetiologies of blepharitis, e.g. allergic contact dermatitis, or infestation with demodex mites, should also be considered in patients with AD. Sometimes, aetiological classification of conjunctivitis is straightforward, e.g. onset of allergic rhinoconjunctivitis during pollen season in AD patients with known allergic disease and positive response to antihistamines, or purulent infectious conjunctivitis in conjunction with impetiginized AD, a positive culture, and a favourable response to antibiotics. At other times, it can be clinically difficult to categorize conjunctivitis and, in this case, more extensive diagnostic workup is required. There was consensus to recommend pursuit of a diagnostic workup for type 1 and 4 allergy in AD patients with conjunctivitis. Notably, in severe conjunctivitis cases, it is important that an ophthalmologist rule out atopic keratoconjunctivitis, which may lead to keratitis and blindness.^{2,19}

There was consensus that dermatologists should prescribe lubricating eye drops, ointments or oral antihistamines in AD patients with conjunctivitis, but also that dermatologists should leave the use of corticosteroid, ciclosporin or tacrolimus eye drops to ophthalmologists. A general concern with corticosteroid eye drop use, besides the risk of superinfection, is induction of increased ocular pressure, which may result in glaucoma. The price of commercially available ciclosporin eyedrop treatment is relatively high and may limit its use, but ciclosporin eye drops are generally well tolerated and are effective for vernal and atopic keratoconjunctivitis.²⁰ Compounding of ciclosporin eye drops in a pharmacy may be an alternative, if available. The use of tacrolimus eye drops (and tacrolimus as ointment) has also proven favourable with no systemic side-effects in a large trial; however, incident herpetic and bacterial infections can occur, warranting ophthalmological monitoring.²¹ Despite the more limited experience with eye drops by dermatologists, rapid access to ophthalmological service may be difficult, sometimes warranting a short course of corticosteroid eye drop without ophthalmological consultations. However, persistent or recurrent conjunctivitis requiring repeated or prolonged use of corticosteroid, tacrolimus and ciclosporin containing eye drops, must be managed by an ophthalmologist, given the risk of glaucoma, cataract and infections.

Conjunctivitis occurring during dupilumab treatment is observed in 9–28% of every 2-week dupilumab users in clinical trials^{4–6} and in real-world evidence from US, Dutch and French registries (oral presentations, EADV, Paris, September, 2018).⁷ Observations in other clinical trials may aid in understanding the pathogenesis of conjunctivitis occurring during dupilumab treatment. For example, the risk of conjunctivitis in patients with asthma receiving treatment with dupilumab is similar to that of controls,²² raising the question of ocular or immune differences between patients with AD and asthma. Furthermore, larger trials of IL-13 inhibitors may indicate the role of IL-13 inhibition directly vs. IL-4 inhibition or the combination through IL4Ra targeting; in small phase II trials, an increased incidence of conjunctivitis was observed with lebrikizumab, but not tralokinumab.^{23,24} One possible mechanism of conjunctivitis occurring during dupilumab treatment involves the observed early increases in blood total eosinophil counts, since eosinophil chemotactic and activating peptides, eotaxin 1 and 2, are increased in the mucus and tears of patients with vernal and atopic keratoconjunctivitis.²⁵ Other hypotheses include a pathogenic role of increased OX40 ligand activity,²⁶ or IL-17 driven inflammation following colonization with *demodex* mites.²⁷ Notably, a recent histopathology case study showed marked depletion of goblet cells in conjunctivae of patients with dupilumab-associated conjunctivitis.²⁸

The limitations of our study include the following: 1) reliance on expert dermatologist opinion; 2) while discussion participants were all dermatologists experienced with dupilumab use

(and managing dupilumab-associated conjunctivitis) from clinical trials and dermatology office use; number of respondents was relatively small; and survey respondents included few councillors from countries lacking access to dupilumab.

Conclusion

This small IEC member survey and round table discussion provided guidance from a group of AD experts regarding general diagnostic workup and management of conjunctivitis occurring during dupilumab treatment. We achieved consensus for several statements regarding management of conjunctivitis in patients with AD, also before and during dupilumab therapy, but emphasize that correct classification and treatment of conjunctivitis is important and that the threshold of contacting ophthalmologists should be low. Topical ophthalmological therapies recommended by IEC members have proved successful and have allowed dupilumab treatment continuation, but consideration of alternative diagnoses that require further assessment and/or initiation of treatment other than topical anti-inflammatory medication is critical before initiation of therapy²⁹; these management decisions may be beyond the ability and comfort of a dermatologist or other dupilumab prescriber and may require joint management with an ophthalmologist. Future studies on the aetiopathogenesis, predictive factors and approach to conjunctivitis occurring during dupilumab treatment are warranted.

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